CERAVE DEVELOPED WITH DERMATOLOGISTS HYDRATING MINERAL SUNSCREEN BROAD SPECTRUM SPF 30 FACE SHEER TINT SUNSCREEN-
titanium dioxide and zinc oxide lotion
L'Oreal USA Products, Inc.

Drug Facts

Active ingredients

Titanium dioxide 5.5%

Zinc oxide 10%

Purpose

Sunscreen

Uses

• helps prevent sunburn
• if used as directed with other sun protection measures (see ***Directions***), decreases the risk of skin cancer and early skin aging caused by the sun

Warnings

For external use only

Do not use

on damaged or broken skin

When using this product

keep out of eyes. Rinse with water to remove.

Stop use and ask a doctor if

rash occurs

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

For sunscreen use:
● apply liberally 15 minutes before sun exposure
● reapply at least every 2 hours
● use a water resistant sunscreen if swimming or sweating
● **Sun Protection Measures**. Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk, regularly use a sunscreen with a Broad Spectrum SPF value of 15 or higher and other sun protection measures including:
● limit time in the sun, especially from 10 a.m. – 2 p.m.
● wear long-sleeved shirts, pants, hats, and sunglasses
● children under 6 months of age: Ask a doctor

Inactive ingredients

water, c12-15 alkyl benzoate, isohexadecane, isononyl isononanoate, dicaprylyl ether, PEG-30 dipolyhydroxystearate, triethylhexanoin, polyglyceryl-4 isostearate, dicaprylyl carbonate, ethylene/acrylic acid copolymer, triethanolamine, silica, poly c10-30 alkyl acrylate, stearic acid, ceramide NP, ceramide AP, ceramide EOP, carbomer, niacinamide, cetearyl alcohol, triethoxycaprylylsilane, behentrimonium methosulfate, sodium chloride, salicylic acid, sodium hyaluronate, sodium lauroyl lactylate, cholesterol, aluminum stearate, alumina, aluminum hydroxide, iron oxides, phenoxyethanol, p-anisic acid, chlorphenesin, tocoferol, disodium EDTA, disodium stearoyl glutamate, propylene carbonate, citric acid, caprylyl glycol, capryloyl salicylic acid, caprylic/capric triglyceride, diethylhexyl syringylidenemalonate, disteardimonium hectorite, xanthan gum, phytosphingosine, polyhydroxystearic acid, ethylhexylglycerin

Other information

protect the product in this container from excessive heat and direct sun

Questions or comments?

Toll-free number **1-888-768-2915**

www.CeraVe.com

Small Sunscreen

Glue

Drug Facts

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Zinc oxide 10%	Sunscreen

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Questions or comments?

Toll-free number **1-888-768-2915**

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No Varnish

CeraVe®
DEVELOPED WITH DERMATOLOGISTS

MINERAL SUNSCREEN

Hydrating Sunscreen
BROAD SPECTRUM SPF 30

30 FACE SHEER TINT

RECOMMENDED SKIN CANCER FOUNDATION DAILY USE

CeraVe® LLC,
New York, NY 10001
Made in USA of US and/or Imported Ingredients
www.cerave.com

3612622240794

CeraVe®
DEVELOPED WITH DERMATOLOGISTS

Hydrating Mineral Sunscreen

BROAD SPECTRUM SPF 30

30 FACE SHEER TINT

Blends seamlessly into skin with a lightweight, non-greasy feel

Suitable for sensitive skin

3 essential ceramides & niacinamide

MINERAL SUNSCREEN

1.7 FL OZ (50 mL)

CeraVe® Hydrating Mineral Sunscreen Sheer Tint
BROAD SPECTRUM SPF 30

Developed with dermatologists, its formula – with 3 essential ceramides – helps restore and maintain the protective skin barrier.

UVA UVB

Broad Spectrum Protection
Helps protect your skin against harmful UVA and UVB rays

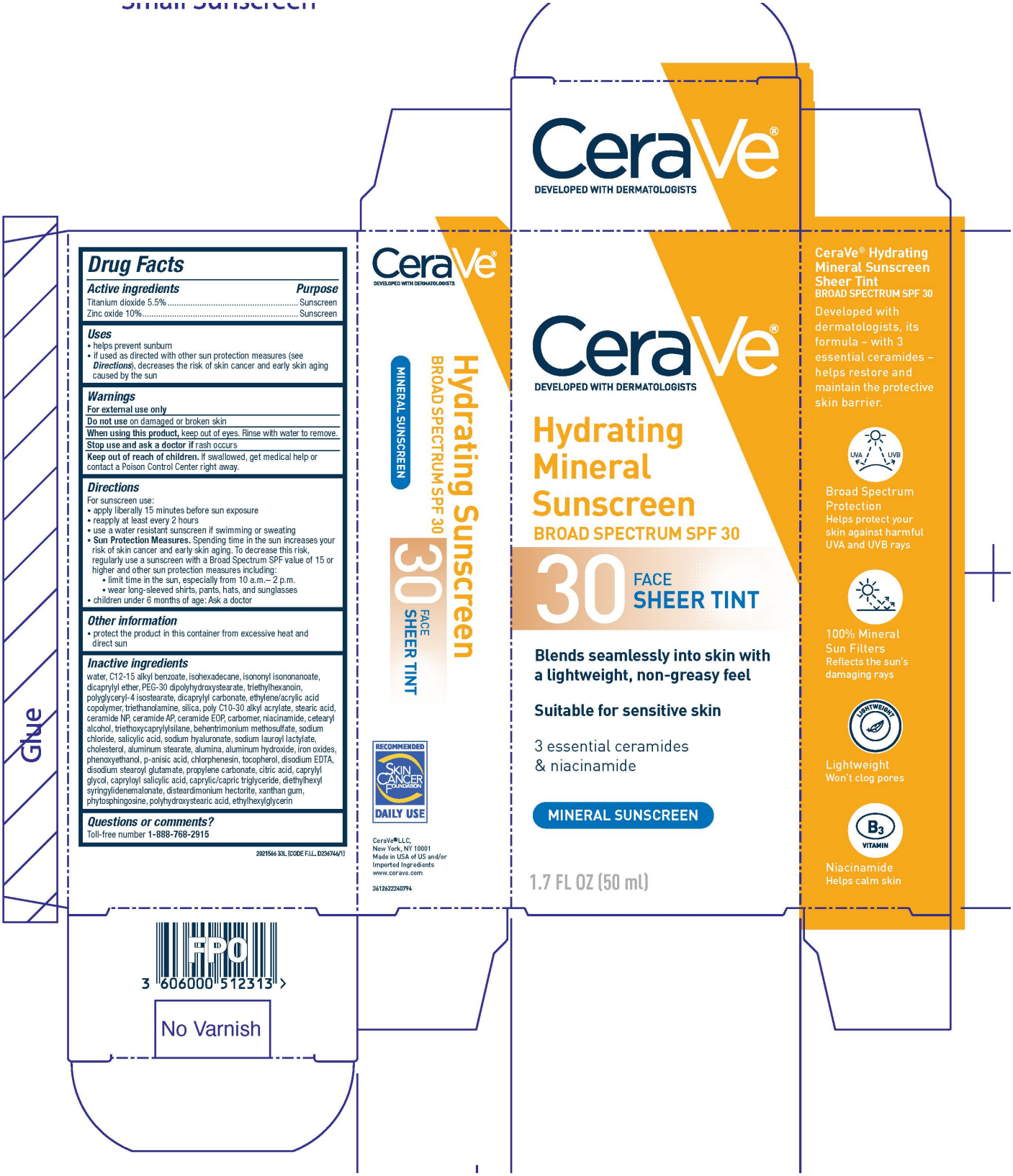
100% Mineral Sun Filters
Reflects the sun's damaging rays

LIGHTWEIGHT

Lightweight
Won't clog pores

B_3 VITAMIN

Niacinamide
Helps calm skin



CERAVE DEVELOPED WITH DERMATOLOGISTS HYDRATING MINERAL SUNSCREEN BROAD SPECTRUM SPF 30 FACE SHEER TINT SUNSCREEN

titanium dioxide and zinc oxide lotion

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49967-231
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	55 mg in 1 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	100 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
ISOHEXADECANE (UNII: 918X1OUF1E)	
ISONONYL ISONONANOATE (UNII: S4V5BS6GCX)	
DICAPRYLYL ETHER (UNII: 77JZM5516Z)	
PEG-30 DIPOLYHYDROXYSTEARATE (UNII: 9713Q0S7FO)	
TRIETHYLHEXANOIN (UNII: 7K3W1BIU6K)	
POLYGLYCERYL-4 ISOSTEARATE (UNII: 820DPX33S7)	
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)	
ACRYLIC ACID/ETHYLENE COPOLYMER (600 MPA.S) (UNII: 1PEZ3NLY6I)	
TROLAMINE (UNII: 9O3K93S3TK)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
CERAMIDE NP (UNII: 4370DF050B)	
CERAMIDE AP (UNII: F1X8L2B00J)	
CERAMIDE 1 (UNII: 5THT33P7X7)	
CARBOMER HOMOPOLYMER TYPE B (ALLYL SUCROSE CROSSLINKED) (UNII: Z135WT9208)	
NIACINAMIDE (UNII: 25X51I8RD4)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
BEHENTRIMONIUM METHOSULFATE (UNII: 5SHP745C61)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
SODIUM LAUROYL LACTYLATE (UNII: 7243K85WFO)	
CHOLESTEROL (UNII: 97C5T2UQ7J)	
ALUMINUM STEARATE (UNII: U6XF9NP8HM)	
ALUMINUM OXIDE (UNII: LMI26O6933)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
P-ANISIC ACID (UNII: 4SB6Y7DMM3)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
TOCOPHEROL (UNII: R0ZB2556P8)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
DISODIUM STEAROYL GLUTAMATE (UNII: 45ASM2L11M)	

PROPYLENE CARBONATE (UNII: 8D08K3S51E)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
CAPRYLOYL SALICYLIC ACID (UNII: 5F7PJF6AA4)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
DIETHYLHEXYL SYRINGYLIDENEMALONATE (UNII: 3V5U97P248)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
XANTHAN GUM (UNII: TTV12P4NEE)	
PHYTOSPHINGOSINE (UNII: GIN46U9Q2Q)	
PHYTOSPHINGOSINE HYDROCHLORIDE (UNII: TT871XV7TU)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49967-231-01	1 in 1 CARTON	12/01/2019	
1		50 mL in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:49967-231-02	1 in 1 CARTON	12/01/2019	
2		5 mL in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	12/01/2019	

Labeler - L'Oreal USA Products, Inc. (002136794)

Establishment

Name	Address	ID/FEI	Business Operations
L'Oreal USA Products, Inc.		624244349	MANUFACTURE(49967-231)

Revised: 12/2023 L'Oreal USA Products, Inc.